agreement. Please note that this agreement limits use to scientific research. Use for software development or design or implementation of commercial gateways or other similar uses is prohibited and may result in loss of user privileges and other penalties.

FILE 'HOME' ENTERED AT 18:06:07 ON 04 JUN 2008

=> fil caplus
COST IN U.S. DOLLARS

COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION FULL ESTIMATED COST 0.63 0.63

FILE 'CAPLUS' ENTERED AT 18:07:52 ON 04 JUN 2008
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.
COPYRIGHT (C) 2008 AMERICAN CHEMICAL SOCIETY (ACS)

Copyright of the articles to which records in this database refer is held by the publishers listed in the PUBLISHER (PB) field (available for records published or updated in Chemical Abstracts after December 26, 1996), unless otherwise indicated in the original publications. The CA Lexicon is the copyrighted intellectual property of the American Chemical Society and is provided to assist you in searching databases on STN. Any dissemination, distribution, copying, or storing of this information, without the prior written consent of CAS, is strictly prohibited.

FILE COVERS 1907 - 4 Jun 2008 VOL 148 ISS 23 FILE LAST UPDATED: 3 Jun 2008 (20080603/ED)

Effective October 17, 2005, revised CAS Information Use Policies apply. They are available for your review at:

http://www.cas.org/legal/infopolicy.html

=> d his

(FILE 'HOME' ENTERED AT 18:06:07 ON 04 JUN 2008)

FILE 'CAPLUS' ENTERED AT 18:07:52 ON 04 JUN 2008

=> s alpha chromium oxide

-1767788 ALPHA

396562 CHROMIUM

1870232 OXIDE

L1 26 ALPHA CHROMIUM OXIDE

(ALPHA(W)CHROMIUM(W)OXIDE)

=> s l1 and crystalline

82961 CRYSTALLINE

L2 2 L1 AND CRYSTALLINE

=> d 1-2 bib abs

L2 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2008 ACS on STN

AN 1977:182159 CAPLUS

DN 86:182159

OREF 86:28487a,28490a

TI On the preparation of finely dispersed crystalline .

alpha.-chromium oxide (α -Cr2O3) Proinova-Mekhandzhieva, R.; Mekhandzhiev, D. ΑU Inst. Chem. Technol., Sofia, Bulg. CS Doklady Bolgarskoi Akademii Nauk (1976), 29(12), 1787-90 SO CODEN: DBANAD; ISSN: 0366-8681 Journal DT LA English Samples of Cr(OH)3 with large sp. surface, obtained by precipitation from AΒ Cr(NO3)3 solns. with NH3 in presence of EtOH, were heated at 200, 300, 400, and 500° with continuous blowing of N, O, and H into the reaction space or in vacuum to produce finely dispersed crystalline α -Cr203. Even at 500° the conversion of Cr(OH)3 into Cr2O3 is not complete. The products obtained were characterized by magnetic susceptibility and sp. surface area detns. and x-ray diffraction. The sp. surface areas of the products obtained at different temps. in the different atms. and in vacuum are given. The samples obtained at 400° in the presence of N had a large sp. surface (160 m2/g) and was antiferromagnetic, i.e., it had the crystal structure of α -Cr203. L2 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2008 ACS on STN ΑN 1963:39547 CAPLUS 58:39547 DN OREF 58:6678e-f Mechanism for the dehydrogenation of cyclohexane on crystalline α -chromium oxide Balandin, A. A.; Rozhdestvenskaya, I. D. ΑU SO Izvestiva Akademii Nauk SSSR, Seriya Khimicheskaya (1961) 1955-60 CODEN: IASKA6; ISSN: 0002-3353 DT Journal LA Unavailable The dehydrogenation of cyclohexane was studied at 330-400° on a AR crystalline α -Cr203 catalyst. The preparation of the catalyst, materials, and the method was described earlier (CA 56, 13594f). A comparison of the exptl. data with the theories for the mechanism of the reaction showed that the data agreed best with the planar sextet mechanism (CA 27,265; 51, 10208f). 34 references. => s l1 and nickel atoms 673165 NICKEL 559825 ATOMS 565 NICKEL ATOMS (NICKEL (W) ATOMS) T.3 1 L1 AND NICKEL ATOMS => d bib abs ANSWER 1 OF 1 CAPLUS COPYRIGHT 2008 ACS on STN L3 2004:182755 CAPLUS ΑN DN 140:219737 ΤТ Nickel-substituted and mixed nickel-and-cobalt-substituted chromium oxide compositions, their preparation, and their use as catalysts and catalyst precursors Nappa, Mario J.; Rao, Velliyur Nott Mallikarjuna; Rosenfeld, H. David; INSubramoney, Shekhar; Subramanian, Munirpallam A.; Sievert, Allen C. PAE.I. du Pont de Nemours and Company, USA SO PCT Int. Appl., 55 pp. CODEN: PIXXD2

Patent

FAN.CNT 1

English

DT LA

```
PATENT NO.
                    KIND DATE
                                        APPLICATION NO. DATE
    WO 2004018095 A1 20040304 WO 2003-US26327 20030821
PΤ
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM,
             PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN,
             TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,
             FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR,
             BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
                              20040311 AU 2003-265592 20030821
20050713 EP 2003-793282 20030821
     AU 2003265592
                         A1
     EP 1551551
                          A1
         R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK
                     A 20051005 CN 2003-819939 20030821
T 20051202 JP 2004-529844 20030821
     CN 1678391
     JP 2005536425
                                          RU 2005-107802
US 2005-523226
RU 2318595 C2 20080310
US 20050227865 A1 20051013
PRAI US 2002-405221P P 20020822
WO 2003-US26327 W 20030821
                                                                    20030821
                                                                    20050131
OS
     CASREACT 140:219737
     A crystalline \alpha -chromium oxide where 0.05-2
AΒ
     atom % of the chromium atoms in the \alpha -chromium
     oxide lattice are substituted by nickel atoms,
     and optionally, addnl. chromium atoms in the alpha-
     chromium oxide lattice are substituted by trivalent
     cobalt atoms (provided that the total amount of the nickel
     atoms and the trivalent cobalt atoms in the \alpha -
     chromium oxide lattice is no more than 6 atom%) is
     disclosed. Also disclosed is a chromium-containing catalyst composition
comprising
     as a chromium-containing component the crystalline substituted lpha -
     chromium oxide; and a method for preparing a composition
     comprising the crystalline substituted \alpha -chromium
     oxide. The method comprises (a) co-precipitating a solid by adding
     ammonium hydroxide to an aqueous solution of a soluble divalent nickel salt, a
     trivalent chromium salt, and optionally, a soluble divalent or trivalent
     cobalt salt, that contains at least three moles of nitrate per mol of
     chromium in the solution, has a nickel concentration 0.05-2 mol% of the total
\circf
     nickel, chromium, and cobalt in the solution, and has a combined concentration
\circf
     nickel and cobalt of no more than 6 mol% of the total of nickel, chromium,
     and cobalt in the solution; and after at least 3 mol of ammonium per mol of
     chromium has been added to the solution; (b) collecting the co-precipitated
solid
     formed in (a); (c) drying the collected solid; and (d) calcining the dried
     solid. Also disclosed is a chromium-containing catalyst composition
comprising a
     chromium-containing component prepared by treating said crystalline substituted
     -chromium oxide with a fluorinating agent; and a process for changing the
     fluorine distribution (i.e., content and/or arrangement) in a hydrocarbon
     or halogenated hydrocarbon in the presence of a catalyst. The process
     involves using as the catalyst a composition comprising the crystalline
substituted
     alpha-chromium oxide and/or the treated
     substituted \boldsymbol{\alpha} -chromium oxide.
             THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 6
```

ALL CITATIONS AVAILABLE IN THE RE FORMAT

```
=> s chromium oxide and nickel atoms
        396562 CHROMIUM
       1870232 OXIDE
         37001 CHROMIUM OXIDE
                 (CHROMIUM(W)OXIDE)
        673165 NICKEL
        559825 ATOMS
           565 NICKEL ATOMS
                 (NICKEL (W) ATOMS)
L4
             2 CHROMIUM OXIDE AND NICKEL ATOMS
=> d 1-2 bib abs
    ANSWER 1 OF 2 CAPLUS COPYRIGHT 2008 ACS on STN
T.4
     2007:1256306 CAPLUS
ΑN
    Metal dusting of nickel and its alloys
ТΤ
ΑU
     Zhang, Jianqiang; Young, David J.
CS
     School of Materials Science and Engineering, University of New South
     Wales, Sydney, NSW, 2052, Australia
     ECS Transactions (2007), 3(14, High Temperature Corrosion and Materials
SO
     Chemistry 6), 27-41
     CODEN: ECSTF8
    Electrochemical Society
PΒ
    Journal; (computer optical disk)
DT
LA
     English
AΒ
     Pure nickel, Ni-Cu binary alloys, Types 304 and 310 stainless steels and
     Alloy 800H were all reacted with carbon-supersatd. CO/H2/H2O at
     680°. In general, the reaction led to growth of an external
     deposit, together with graphitization at both external surfaces and
     interior grain boundaries. By independently controlling the supersatd.
     carbon activity and pCO, the reaction of pure nickel was determined by
     parallel, independent reaction paths, and the carbon uptake rate was well
     described by Rate = k1pCOpH2 + k2p2CO - k3p2H2 leading to a maximum rate at
     about pCO .apprx. 0.7 atmospheric The surface graphite layer thickened,
growing
     into the metal, while graphite particle clusters and nanofilaments formed
     on the surface. Each particle cluster and filament contained metal
     nanoparticles. Addition of copper to the nickel suppressed graphite particle
     cluster formation, thereby decreasing greatly the overall dusting rate.
     This phenomenon is interpreted in terms of graphite nucleation and copper
     interference with nucleation sites involving multiple nickel
     atoms. Reaction conditions used were such as to stabilize Fe3C
     and chromium carbides, Cr203 and spinel, but not iron or nickel oxides.
     Under these conditions, all three alloys, 304SS, 310SS and 800H, dusted,
     with 304SS experiencing the greatest metal wastage. Addns. of copper had
    no effect on the dusting rate of 304SS, but greatly decreased the dusting
     of 310SS and 800H.
RE.CNT 31
              THERE ARE 31 CITED REFERENCES AVAILABLE FOR THIS RECORD
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
L4
     ANSWER 2 OF 2 CAPLUS COPYRIGHT 2008 ACS on STN
ΑN
     2004:182755 CAPLUS
DN
     140:219737
ΤI
     Nickel-substituted and mixed nickel-and-cobalt-substituted
     chromium oxide compositions, their preparation, and
     their use as catalysts and catalyst precursors
    Nappa, Mario J.; Rao, Velliyur Nott Mallikarjuna; Rosenfeld, H. David;
ΙN
```

Subramoney, Shekhar; Subramanian, Munirpallam A.; Sievert, Allen C.

E.I. du Pont de Nemours and Company, USA

PA

```
SO
     PCT Int. Appl., 55 pp.
     CODEN: PIXXD2
DТ
     Patent
LA
     English
FAN.CNT 1
     WO 2004018095 A1
     PATENT NO.
                        KIND DATE
                                         APPLICATION NO.
                                                                     DATE
                         A1 20040304 WO 2003-US26327 20030821
                                                                     -----
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM,
             PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN,
             TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,
             FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR,
             BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
     AU 2003265592 A1 20040311 AU 2003-265592 20030821
EP 1551551 A1 20050713 EP 2003-793282 20030821
         R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK
                     T, Lv,

A 200

T 20051202

C2 20080310

A1 20051013

P 20020822

T 20030821
                                           CN 2003-819939
JP 2004-529844
RU 2005-107802
US 2005-523226
                                                                     20030821
     JP 2005536425
     RU 2318595
                                                                      20030821
     US 20050227865
                                                                      20050131
PRAI US 2002-405221P P
     WO 2003-US26327
OS
     CASREACT 140:219737
     A crystalline \alpha- chromium oxide where 0.05-2 atom % of
AΒ
     the chromium atoms in the \alpha- chromium oxide
     lattice are substituted by nickel atoms, and
     optionally, addnl. chromium atoms in the alpha-chromium
     oxide lattice are substituted by trivalent cobalt atoms (provided
     that the total amount of the nickel atoms and the
     trivalent cobalt atoms in the \alpha- chromium oxide
     lattice is no more than 6 atom%) is disclosed. Also disclosed is a
     chromium-containing catalyst composition comprising as a chromium-containing
     the crystalline substituted \alpha- chromium oxide; and a
     method for preparing a composition comprising the crystalline substituted \alpha-
     chromium oxide. The method comprises (a) co-precipitating a
     solid by adding ammonium hydroxide to an aqueous solution of a soluble divalent
     nickel salt, a soluble trivalent chromium salt, and optionally, a soluble
     divalent or trivalent cobalt salt, that contains at least three moles of
     nitrate per mol of chromium in the solution, has a nickel concentration 0.05-2
mol%
     of the total of nickel, chromium, and cobalt in the solution, and has a
     combined concentration of nickel and cobalt of no more than 6 mol% of the total
     of nickel, chromium, and cobalt in the solution; and after at least 3 mol of
     ammonium per mol of chromium has been added to the solution; (b) collecting
     the co-precipitated solid formed in (a); (c) drying the collected solid; and
(d)
     calcining the dried solid. Also disclosed is a chromium-containing catalyst
     composition comprising a chromium-containing component prepared by treating
said
     crystalline substituted -chromium oxide with a fluorinating
     agent; and a process for changing the fluorine distribution (i.e., content
     and/or arrangement) in a hydrocarbon or halogenated hydrocarbon in the
     presence of a catalyst. The process involves using as the catalyst a
     composition comprising the crystalline substituted alpha-chromium
```

oxide and/or the treated substituted $\alpha\text{--}$ chromium

oxide.

RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

=>

---Logging off of STN---

=>

Executing the logoff script...

=> LOG Y

COST IN U.S. DOLLARS	SINCE FILE	TOTAL
	ENTRY	SESSION
FULL ESTIMATED COST	39.11	39.74
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	-4.00	-4.00

STN INTERNATIONAL LOGOFF AT 18:11:46 ON 04 JUN 2008